



CERTIFICATION

I, Takashi KOJIMA of Ginza Ohtsuka Bldg., 2F, 16-12, Ginza 2-chome, Chuo-ku, Tokyo, Japan, hereby certify that I am the translator of the accompanying certified official copy of the documents in respect of an application for a patent filed in Japan on the 6th of July, 2000 and of the official certificate attached thereto, and certify that the following is a true and correct translation to the best of my knowledge and belief.

Dated this 12th day of May, 2003

Takashi KOJIMA

(Translation)

PATENT OFFICE JAPANESE GOVERNMENT

This is to certify that the annexed is a true copy of the following application as filed with this Office.

Date of Application: July 6, 2000

Application Number: Japanese Patent Application

No. 2000-204728

Applicant(s): Nisshinbo Industries, Inc.

March 23, 2001

Commissioner,

Patent Office

Kozo OIKAWA

(sealed)

2000-204728

[SPECIFICATION]

[TITLE OF THE INVENTION] Fuel Cell Separator, Process for Production Thereof, and Polymer Electrolyte Fuel Cell

[CLAIMS]

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[Claim 1] A fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein the electrically conductive carbon powder is present such that its particles longer than 50 μm at maximum in the major axis direction and longer than 30 μm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional area in the vertical direction.

[Claim 2] A fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein said binding agent is contained in an amount of 10 to 50 parts by mass for 100 parts by mass of the electrically conductive carbon powder and the electrically conductive carbon powder is spherical or massive graphite having a mean particle diameter of 100 to 500 $\mu m.$

[Claim 3] The fuel cell separator as defined in Claim 2, wherein the spherical or massive graphite has a bulk density higher than 0.6~g/ml.

[Claim 4] The fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent as defined in Claim 2 or 3, wherein the electrically conductive carbon powder is present such that its particles longer than 50 μm at maximum in the major axis direction and longer than 30 μm

at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional area in the vertical direction.

[Claim 5] The fuel cell separator as defined in any one of Claims 1 to 4, which has a resistivity not higher than 20 m Ω cm measured according to JIS H0602.

[Claim 6] A process for producing a fuel cell separator having gas supply grooves on one side or both sides thereof from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein said process comprising injection molding a mixture containing 10 to 50 parts by mass of a binding agent for 100 pars by mass of the electrically conductive carbon powder.

[Claim 7] The fuel cell separator as defined in Claim 6, wherein the electrically conductive carbon powder is spherical or massive graphite having a mean particle diameter of 100 to 500 μm , and the spherical or massive graphite having a bulk density higher than 0.6 g/ml is used.

[Claim 8] A polymer electrolyte fuel cell consisting of a plurality of unit cells connected together, each unit cell consisting of a pair of electrodes holding a polymer electrolyte membrane between them and a pair of separators holding the electrodes between them, said separator having passages molded thereon through which gas is supplied and discharged, characterized in that all or part of the separators in the fuel cells are those which are defined in any one of Claims 1 to 5.

[DETAILED DESCRIPTION OF THE INVENTION]

[0001]

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[Technical Field of the Invention]

The present invention relates to a fuel cell separator, a process for production thereof, and a polymer electrolyte fuel cell. More particularly, the present invention relates to a fuel cell separator having high elasticity, high electrical conductivity, and good moldability and a process for production thereof. The

present invention relates also to a fuel cell having good gas sealing performance and good impact resistance in which all or part of the separators are those which are mentioned above. The fuel cell is suitable for use as a mobile power source for cars, hybrid cars, and small ships.

[0002]

[Prior Art and the Problem to be Solved by the Invention]
A polymer electrolyte fuel cell is composed of tens to hundreds of fuel cells (as unit cells) connected together.
Each fuel cell consists of two fuel cell separators 1, 1 and one polymer electrolyte membrane 2 and two gas diffusion electrodes 3 which are held between the separators, each separator having a plurality of ribs 1a on both sides thereof, as shown in Fig. 1.

[0003]

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The above-mentioned fuel cell separator 1 is a thin platy body having a plurality of ribs 1a on both sides thereof and a plurality of gas feed grooves 4 on one side or both sides thereof, as shown in Figs. 2A and 2B. The ribs 1a of the separator and the electrode 3 form passages 4 for fuel gas such as hydrogen and oxygen to be supplied and discharged. Therefore, the fuel cell separators are required to have high elasticity and good dimensional accuracy. Moreover, the fuel cell separators and unit fuel cells are required to have good gas seal performance to prevent the leakage of fuel gas, good resistance to cracking by tightening at the time of assembling, and good impact resistance for the fuel cell to be used as a mobile power source for automobiles.

30 [0004]

To meet these requirements, there has been proposed a separator for polymer electrolyte fuel cell in Japanese Patent Laid-open No. Hei 11-297337. This separator is obtained by curing from a homogenous mixture composed of 100 parts by weight of carbonaceous powder and 10 to 100 parts by weight of thermosetting resin. According to this disclosure, the carbonaceous powder is a graphite powder

having a maximum particle diameter of 125 μm or below. There has also been disclosed a fuel cell separator in Japanese Patent Publication No. 2000-100453. This separator contains expansible graphite particles having a number-average particle diameter of 25 μm or above, preferably 25 to 500 μm . [0005]

Conventional fuel cell separators are made of a highly filled material which encounters difficulties in injection molding. The disadvantage of the above-mentioned separators containing graphite powder is that the fine graphite powder lowers the fluidity of the molding material, adversely affecting injection moldability and mechanical properties. Flake graphite has such a low bulk density that the material containing it slips on the screw of the injection molding machine, making itself incapable of molding. Graphite with a comparatively high bulk density is easily broken into fine powder at the time of mixing or injection molding.

[0006]

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Since a unit cell merely produces a low voltage, it is necessary to connect tens to hundreds of unit cells if a practical output (up to hundreds of kW) is to be obtained. Therefore, there is an urgent demand for a technology that permits efficient mass production of fuel cell separators having a uniform shape free of strain and thickness variation.

[0007]

Conventional fuel cell separators are made of a composition composed of a thermosetting resin such as phenolic resin, and graphite. This composition is incorporated with a large amount of graphite so that the resulting separator has electrically conductivity as required. Therefore, it lacks fluidity and presents difficulties in injection molding. Actual production of separators is by compression molding, which involves heating at 150 to 160°C and pressing at 14.7 to 29.4 MPa for 5 to 10 minutes. Compression molding is slow and inefficient for mass production.

[8000]

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The present invention was completed in view of the foregoing. It is an object of the present invention to provide a high-quality fuel cell separator having high elasticity, high electrical conductivity, and good gas impermeability. It is another object of the present invention to provide a process for producing the fuel cell separator, the process being suitable for efficient, mass production. It is another object of the present invention to provide a polymer electrolyte fuel cell having good gas sealing performance and good impact resistance in which the separators are immune to cracking by tightening at the time of assembling.

[0009]

[Means for Solving the Problem and Embodiment of the Invention]

In order to achieve the above-mentioned object, the present inventors carried out extensive studies. As the result, it was found that a molding composition easily affords a fuel cell separator having good mechanical strength and high electrical conductivity if it contains a large amount of electrically conductive carbon powder of high bulk density.

[0010]

In other words, it was found that the above-mentioned object is achieved by a fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent. According to the present invention, the composition contains 100 parts by mass of the electrically conductive carbon powder and 10 to 50 parts by mass of the binding agent, and the electrically conductive powder is spherical or massive graphite of high bulk density having a mean particle diameter of 100 to 500 μm . The electrically conductive carbon powder is present such that its particles longer than 50 μm at maximum in the

major axis direction and longer than 30 μm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional The fuel cell separator area in the vertical direction. meeting the above-mentioned requirements has high elasticity, high electrical conductivity, and good gas impermeability. A stack of fuel cells in which all or part of the separators in the fuel cells are those which are mentioned above is immune to cracking by tightening at the time of assembling; therefore, it offers good gas seal performance and good impact resistance. It is suitable for use as a portable power source for cars, hybrid cars, and The above-mentioned finding is the basis of small ships. the present invention.

[0011]

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According to the present invention, the fuel cell separator is molded from a mixture (molding composition) containing 100 parts by mass of electrically conductive powder and 10 to 50 parts by mass of binding agent. This molding composition is suitable for injection molding for economical, efficient, mass production of fuel cell separators which are a thin platy body of peculiar shape having gas supply grooves on one side or both sides thereof. [0012]

The present invention is directed to a fuel cell separator, a process for production thereof, and a polymer electrolyte fuel cell, as explained in the following.

The first aspect of the present invention covers a fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein the electrically conductive carbon powder is present such that its particles longer than 50 μm at maximum in the major axis direction and longer than 30 μm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy

more than 50% of the sectional area in the vertical direction.

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The second aspect of the present invention covers a fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein the binding agent is contained in an amount of 10 to 50 parts by mass for 100 parts by mass of the electrically conductive carbon powder and the electrically conductive carbon powder is spherical or massive graphite having a mean particle diameter of 100 to 500 μm .

The third aspect of the present invention covers the fuel cell separator as defined in the second aspect, wherein the spherical or massive graphite has a bulk density higher than $0.6~\rm g/ml$.

The fourth aspect of the present invention covers the fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent as defined in the second or third aspect, wherein the electrically conductive carbon powder is present such that its particles longer than 50 μm at maximum in the major axis direction and longer than 30 μm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional area in the vertical direction.

The fifth aspect of the present invention covers the fuel cell separator as defined in any one of the first to fourth aspects, which has a resistivity not higher than 20 m Ω ·cm measured according to JIS H0602.

The sixth aspect of the present invention covers a process for producing a fuel cell separator having gas supply grooves on one side or both sides thereof from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein the process

comprising injection molding a mixture containing 10 to 50 parts by mass of a binding agent for 100 pars by mass of the electrically conductive carbon.

The seventh aspect of the present invention covers the fuel cell separator as defined in the sixth aspect, wherein the electrically conductive carbon powder is spherical or bulky graphite having a mean particle diameter of 100 to 500 μm , and the spherical or bulky graphite has a bulk density higher than 0.6 g/ml.

The eighth aspect of the present invention covers a polymer electrolyte fuel cell consisting of a plurality of unit cells connected together, each unit cell consisting of a pair of electrodes holding a polymer electrolyte membrane between them and a pair of separators holding the electrodes between them, the separator having passages molded thereon through which gas is supplied and discharged, characterized in that all or part of the separators in the fuel cells are those which are defined in any one of the first to fifth aspects.

[0013]

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The invention will be described in more detail below. According to the present invention, the fuel cell separator which has gas supply grooves on one side or both sides thereof is formed from a composition composed mainly of an electrically conductive carbon powder and a binding agent. The electrically conductive carbon powder is present such that its particles longer than 50 µm at maximum in the major axis direction and longer than 30 µm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional area in the vertical direction. Because of these characteristics properties, the fuel cell separator has good mechanical properties, electrical conductivity, and gas impermeability.

[0014]

The cross section specified above is noticed when the fuel cell is cut vertical in the thickness direction and the cut surface is observed under a microscope provided with a digital CCD camera ("Axioscope 2" made by Carl Zeiss Co., Ltd.). The fuel cell separator is a flat platy body, measuring 120 mm long, 120 mm wide, and 2.3 mm thick, having gas supply grooves on one side or both sides thereof, as shown in Figs. 2A and 2B. An example of the microphotograph (× 200) thus obtained is shown in reference photographs 1 and 2. White irregular shapes represent graphite particles. Incidentally, the magnification should preferably be 50 to 500 times, more preferably 50 to 200 times.

[0015]

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The thus obtained digital microphotograph is examined as follows for irregular graphite particles to estimate their size and their area occupying in the photograph. First, the maximum lengths in the major axis direction and the minor axis direction are measured, and graphite particles are selected which have a maximum length of 50 µm and above in the major axis direction and a maximum length of 30 μm and above in the minor axis direction, as shown in The approximate area of a graphite particle is calculated by multiplying the length in the major axis direction by the length in the minor axis direction. not always necessary that the maximum length in the major axis direction is the length that passes through the center of a graphite particle. Instead, the longest line across a graphite particle is regarded as the maximum length in the major axis direction. The longest line perpendicular to the maximum line in the major axis direction is regarded as the maximum length in the minor axis direction. A hole may be neglected unless it is large enough to split a graphite particle. In the case of a graphite particle having a very complex shape, it is possible to obtain its area by dividing it into several parts.

[0016]

To be more specific, the schematic diagram in Fig. 3 (illustrating the vertical cross section) permits one to calculate the approximate total area (S1) of graphite particles by summing up (a1 \times b1) + (a2 \times b2) + (a3 \times b3) + (a4 \times b4)... The ratio of the area occupied by graphite particles is calculated from (S1/S2) \times 100, where S1 is the total area occupied by graphite particles and S2 is the area of the vertical cross section (x \times y in Fig. 3).

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According to the present invention, the maximum length in the major axis direction in the vertical cross section should be 50 µm or above, preferably 70 µm or above, and the maximum length in the minor axis direction in the vertical cross section should be 30 µm or above, preferably 50 µm or above, and the ratio of the area occupied by graphite particles should be 50% or above, preferably 50 to 80%, more preferably 55 to 75%, and most desirably 60 to 75%. If the ratio of the area occupied by graphite particles is smaller than specified above, the resulting fuel cell separator does not have characteristic properties required. Conversely, if it is larger than specified above, the resulting fuel cell separator has remarkably low mechanical strength.

[0018]

The fuel cell separator according to the present invention is formed from a composition composed mainly of (A) an electrically conductive carbon powder and (B) a binding agent.

[0019]

The electrically conductive carbon powder as component (A) is either natural one or artificial one, such as carbon black, Ketjen black, acetylene black, carbon whisker, and graphite. They may be used alone or in combination with one another. Of these examples, graphite is particularly preferable.

[0020]

The graphite used in the present invention is spherical graphite or massive graphite. Flake graphite is not desirable because it gives such defects as flow marks and weld lines to the molded product, or it causes cracking. In addition, flake graphite has a low bulk density and hence causes the molding composition to slip on the screw of the injection molding machine. The spherical graphite or massive graphite should have a bulk density of 0.6 g/ml or above, preferably 0.7 g/ml or above, more preferably 0.7 to 1.2 g/ml, and most desirably 0.7 to 1.0 g/ml, which is measured according to JIS K6891-1995. With an excessively low bulk density, the graphite causes the above-mentioned troubles and adversely affects the moldability of the molding composition.

[0021]

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The electrically conductive carbon powder should be one which has a mean particle diameter of 100 to 500 μm , preferably 150 to 450 μm , more preferably 150 to 400 μm . One which is coarser than specified above will make the molding composition poor in fluidity and presents difficulties in injection molding. Conversely, one which is finer than specified above will have an adverse effect on the mechanical properties or gas impermeability of the separator.

[0022]

The binding agent as component (B) above is not specifically restricted. It may be selected from any of the following thermosetting resins and thermoplastic resins. Examples of thermosetting resins include phenolic resin such as resol type or novolak type, furan resin such as furfuryl alcohol resin and furfuryl alcohol phenolic resin, polyimide resin, polycarbodiimide resin, polyacrylonitrile resin, pyrene-phenanthrene resin, polyvinyl chloride resin, epoxy resin, urea resin, diallylphthalate resin, unsaturated polyester resin, and melamine resin. They may be used alone or in combination with one another.

[0023]

Examples of thermoplastic resins include styrene resins such as acrylonitrile-butadiene-styrene resin (ABS), acrylonitrile-styrene copolymer (AS), high-impact polystyrene (HIPS), polystyrene (PS), methyl methacrylatebutadiene-styrene copolymer (MBS), methyl methacrylatestyrene copolymer (MS), acrylonitrile-ethylene-propylene rubber-styrene copolymer (AES), and acrylonitrile-styreneacrylate (AAS), polyolefin resins such as polyethylene (PE), polypropylene (PP), polybutene-1, ethylene-vinyl acetate 10 copolymer (EVA), and ethylene-vinyl alcohol copolymer (EVOH), polyamide resin, thermoplastic polyester resin, polycarbonate resin (PC), wholly aromatic polyester resin, polyphonylene sulfide (PPS), polyvinyl chloride resin (PVC), polysulfone resin, polyether-ether-ketone resin, (modified) 15 polyphenylene ether resin, polyoxymethylene (POM), polymethyl methacrylate (PMMA), fluoroplastic resin, polyketone resin (PK), norbornane, polyamideimide (PAI), and polyphthalamide (PPA). They may be used alone or in combination with one another. 20

[0024]

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The binding agent as component (B) should be used in an amount of 10 to 50 parts by mass, preferably 10 to 40 parts by mass, more preferably 15 to 30 parts by mass, for 100 parts by mass of the electrically conductive carbon powder as component (A). If the amount of the binding agent as component (B) is less than specified above, the resulting molding composition is poor in fluidity and presents difficulties in injection molding. Conversely, if it is more than specified above, the molding compound cannot be incorporated sufficiently with the electrically conductive carbon powder and hence the resulting separator is poor in electrical conductivity.

[0025]

The molding composition of the fuel cell separator of the present invention may be incorporated with, in addition to the above-mentioned components (A) and (B), optional additives such as fibrous base material, mold release, metal powder, and hydrolysis resisting agent, for improvement in strength, mold releasability, hydrolysis resistance, and electrical conductivity.

[0026]

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The above-mentioned fibrous base material includes inorganic fiber and organic fiber. Inorganic fiber includes metal fiber such as iron, copper, brass, bronze, aluminum or the like, ceramics fiber, potassium titanate fiber, glass fiber, carbon fiber, rock wool, wollastonite, sepiolite, attapulgite, and artificial mineral fiber. Organic fiber includes aramid fiber, polyimide fiber, polyamide fiber, phenolic fiber, cellulose, and acrylic fiber. They may be used alone or in combination with one another. The fibrous base material should be used in an amount of 0 to 10 parts by mass for 100 parts by mass of the electrically conductive carbon powder as component (A).

[0027]

The above-mentioned mold release is not specifically restricted; it includes silicone-based ones, fluorine-based one, metallic soap-based ones, amide-based ones, and wax-based ones. Internal mold release agents such as carnauba wax, stearic acid, and montanic acid are preferable. The mold release should be used in an amount of 0 to 3 parts by mass for 100 parts by mass of the electrically conductive carbon powder as component (A).

[0028]

The above-mentioned metal powder includes those of stainless steel, gold, silver, copper, platinum, titanium, aluminum, and nickel. The metal powder should be one which has a mean particle diameter of 5 to 30 $\mu m\,.$

[0029]

According to the present invention, the fuel cell separator having gas supply grooves on one side or both sides thereof is produced by injection molding from a mixture of 100 parts by mass the electrically conductive carbon powder as component (A) and 10 to 50 parts by mass of

the binding agent as component (B). It is desirable that the electrically conductive carbon powder be spherical or massive graphite having a mean particle diameter of 100 to 500 μ m and a bulk density of 0.6 g/ml or above.

[0030]

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The above-mentioned components for the molding composition should be mixed by means of single-screw extruder or twin-screw extruder or general-purpose mixer. The resulting mixture may be pelletized or merely dried. Mixing by means of a mixer is desirable from the standpoint of not breaking the particles of the electrically conductive carbon powder.

[0031]

The thus obtained mixture undergoes injection molding. The conditions of injection molding vary depending on the injection molding machine and the kind and amount of the binding agent. The following conditions are usually desirable.

Cylinder temperature: 50 to 100°C (at front)

Injection pressure: 8 to 20 MPa

Injection time: 5 to 15 seconds

Mold temperature: 140 to 200°C

Curing time: 15 to 200 seconds

[0032]

Injection molding is not an only method of producing the fuel cell separator of the present invention. Other known molding methods that can be used individually or in combination include compression molding, injection-compression molding, transfer molding, extrusion molding, hydrostatic molding, belt pressing, and roll molding.

[0033]

Owing to the electrically conductive carbon powder having a high bulk density, the production by injection molding according to the present invention permits stable injection molding with complete filling into the mold. Injection molding permits the production of the fuel cell separator of unique shape having gas supply grooves on one

side or both sides thereof, as shown in Figs. 2A and 2B. Therefore, it is suitable for efficient mass production. [0034]

The fuel cell separator obtained as mentioned above should have a resistivity not higher than 20 m Ω ·cm, preferably not higher than 15 m Ω ·cm, more preferably 2 to 15 m Ω ·cm, which is measured according to JIS H0602 providing the four point probe method for measuring the resistivity of silicon single crystal and silicon wafer.

10 [0035]

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The fuel cell separator of the present invention should have mechanical strength such that a test piece (100 \times 10 \times 4 mm) prepared from the molding composition according to JIS K6911 proving "General test methods for thermosetting plastics" has a flexural strength of 20 to 80 MPa, preferably 25 to 60 MPa, a flexural modulus of 1 to 15 GPa, preferably 2 to 10 GPa, and a strain of 2 to 15 mm, preferably 3 to 12 mm.

[0036]

The fuel cell separator of the present invention should have gas permeability such that a specimen (2 mm thick, 100 mm in diameter) prepared from the molding composition according to JIS K7126 providing "Method of evaluating the gas permeability of plastics film" (method B, equal pressure method) has a nitrogen gas permeability at 23°C not higher than 20 ml/m²·24 hr·atm, preferably 2 to 20 ml/m²·24 hr·atm, and more preferably 5 to 15 ml/m²·24 hr·atm. [0037]

The present invention covers a polymer electrolyte fuel cell consisting of a plurality of unit cells connected together, each unit cell consisting of a pair of electrodes holding a polymer electrolyte membrane between them and a pair of separators holding the electrode between them, the separator having passages molded thereon through which gas is supplied and discharged. This polymer electrolyte fuel

cell is characterized in that all or part of the separators in the unit cells are those of the present invention.
[0038]

The stack of fuel cells consists of unit cells, each unit cell consisting of a fuel cell and a pair of separators, as shown in Fig. 1. The fuel cell consists of one polymer electrolyte membrane 2 and an electrode 3 for fuel gas and an electrode 3 for oxidizing agent tightly holding the membrane between them. The separators pertain to the present invention. One separator 1, 1 adjacent to the electrode for fuel gas has a plurality of passages 4 and manifolds for fuel gas, and the other separator 1 adjacent to the electrode for oxidizing agent has a plurality of passages 4 and manifolds for oxidizing agent.

[0039]

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These unit cells are stuck in series and tightened between insulating plates supported by pressing plates. One electrode is supplied with fuel gas and the other electrode is supplied with oxidizing agent gas, so that the stack of fuel cells generates direct current.

[0040]

According to the present invention, the stack of fuel cells is characterized in that all or part of separators therein are those which have high elasticity and good gas impermeability as defined above in the present invention. To be more specific, the percentage of the separators of the present invention in all the separators of the stack of fuel cells should be higher than 50%, preferably 50 to 100%, more preferably 70 to 100%, and most desirably 80 to 100%. If this percentage is lower than specified above, the stack of fuel cells will be poor in gas seal performance and impact resistance because the separators are liable to cracking by tightening at the time of assembling. In the case where the separators of the present invention are used partly, the rest may be supplied with conventional separators.

[0041]

The above-mentioned polymer electrolyte membrane may be any ordinary one which is used for polymer electrolyte fuel cell. For example, it may be a proton-conducting ion-exchange membrane made of fluoroplastic resin, such as polytrifluorosulfonic acid and perfluorocarbonsulfonic acid ("Nafion"). The electrolyte membrane is coated with "Nafion 117 solution", which is a paste of a catalyst dispersed in a mixed solvent of water and lower aliphatic alcohol containing perfluorocarbon sulfonic acid. The catalyst is platinum or platinum alloy supported on carbon powder.

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The pair of electrodes holding the polymer electrolyte membrane between them may be formed from carbon paper, carbon felt, or carbon cloth woven from carbon fiber.

[0043]

The two electrodes are bonded to both sides of the polymer electrolyte membrane so that they are made into one piece. Bonding may be accomplished by hot-pressing at 120 to 130°C or with the aid of an adhesive.

[0044]

The unified combination of the electrodes and electrolyte membrane is held between a pair of separators in such a way that the separators form passages for fuel gas supply and discharge. Thus there is obtained the unit cell. Firm bonding may be ensured with an adhesive applied to the ribs of the separator in contact with the electrode.

[0045]

The polymer electrolyte fuel cell which is constructed according to the present invention is immune to the cracking of separators at the time of assembling, exhibits good gas sealing performance, and high impact resistance. These advantages stem from that fact that all or part (preferably more than 50%) of the separators in the fuel cell are those separators pertaining to the present invention which have high elasticity and good gas impermeability. By virtue of these characteristic properties, the fuel cell is suitable

for use as a mobile power source for cars, hybrid cars, and small ships.

[0046]

Incidentally, the polymer electrolyte fuel cell according to the present invention will find use not only as a portable power source for cars, hybrid cars, small ships, etc. but also as a power source for local electric supply, domestic use, camping sites, artificial satellites, and space development.

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[Effect of the Invention]

The present invention permits efficient mass production of fuel cell separators having high elasticity, high electrical conductivity, and good gas impermeability. The polymer electrolyte fuel cell, in which all or part of separators are those pertaining to the present invention, is immune to the cracking of separators at the time of assembling, and exhibits good gas sealing performance and high impact resistance.

20 [0048]

[EXAMPLES]

The invention will be described in more detail with reference to the following examples and comparative examples, which are not intended to limit the scope thereof. In Table 1, the amount of each component is given in terms of parts by mass.

[0049]

[Examples 1 and 2 and Comparative Examples 1 to 4]

The components in each example shown in Table 1 were mixed together. After drying, the resulting mixture was injection-molded into a fuel cell separator under the following conditions by using an injection molding machine (100F-45K, made by Matsuda Seisakusho). The fuel cell separator measures 120 mm long, 120 mm wide, and 2.3 mm thick, and has gas supply grooves 4 on both sides thereof as shown in Fig. 2A. Incidentally, in Comparative Examples 1 and 4, the molding composition was unable to be fed into the

cylinder or transferred through the cylinder because of slipping on the screw.

<Molding conditions>

Cylinder temperature: 65°C (at front)
Injection pressure: 10 to 15 MPa

Injection time: 5 to 15 seconds

Mold temperature: 160 to 165°C

Curing time: 90 to 120 seconds

[0050]

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The thus obtained separator was examined for moldability, gas impermeability, and the ratio of area occupied by graphite particles in the vertical cross section. The results are shown in Table 1. Reference photograph 1 shows the vertical cross section (x 200) of the separator in Example 2. Reference photograph 2 shows the vertical cross section (x 200) of the separator in Comparative Example 2.

Moldability

O: good

△ : fair

 \times : poor

Gas impermeability

A specimen (2 mm thick, 100 mm in diameter) cut out of the separator was tested for N_2 gas permeability (ml/m²·24 hr·atm) at 23°C according to JIS K7126 providing "Method of evaluating the gas permeability of plastics film" (method B, equal pressure method). The results are rated according to the following criterion.

O: less than 20

 \triangle : 20 to 10^3

 \times : more than 10 3

Ratio of area occupied by graphite particles

An arbitrary vertical cross section (cut in the thickness direction) of the separator was observed under a microscope (\times 200) provided with a digital CCD camera

("Axioscope 2" made by Carl Zeiss Co., Ltd.) for calculation of the ratio of area occupied by graphite particles having the maximum length larger than 50 μ m in the major axis direction and the maximum length larger than 30 μ m in the minor axis direction. No data were available in Comparative Examples 1 and 4 because the molding composition was incapable of injection molding. Data in Comparative Examples 2 and 3 are inaccurate because the resulting separators were not uniform in properties.

10 [0051]

Specimens (each measuring $100 \times 100 \times 4$ mm) were molded from the molding compositions shown in Table 1. They were tested for flexural strength, flexural modulus, strain, and resistivity. The results are shown in Table 1.

15 Flexural strength, flexural modulus, and strain

Measured according to JIS K6911 providing the general test methods for thermosetting plastics.

Resistivity

Measured according to JIS H0602 (four point probe method for silicon single crystal and silicon wafer).

[0052] [Table 1]

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	Example		Comparative Example			
	1	2	1	2	3	4
Graphite A	100	-	_	-	-	4
Graphite B	-	100	ı	-	-	
Graphite C	-	-	100	-	-	-
Graphite D	_	_	-	100	-	•
Graphite E	-	_	_	-	100	-
Graphite F	-	-	-	-	_	100
Phenolic resin *1	17.6	17.6	17.6	17.6	17.6	17.6
Mold release *2	1.2	1.2	1.2	1.2	1.2	1.2
Ratio of area occupied by graphite particles (%)	71	65	_	62	_	-
Gas impermeability	0	0	-	×	×	-
Moldability	0	0	×	Δ	Δ	×
Resistivity (mΩ·cm)	12	13		18	20	
Flexural strength (MPa)	42	46	_	39	50	-
Flexural modulus (GPa)	42	40	_	43	49	-
Strain (mm)	0.6	0.6	-	0.7	0.6	-

Graphite A: massive, having a mean particle diameter of 250 to 425 $\,\mu m$ and a bulk density of 0.9 g/ml.

Graphite B: massive, having a mean particle diameter of 180 to 250 $\,$ μm and a bulk density of 0.8 g/ml.

10 Graphite C: massive, having a mean particle diameter of 60 μm and a bulk density of 0.58 g/ml.

Graphite D: flake, having a mean particle diameter of 425 μm and a bulk density of 0.5 g/ml.

Graphite E: flake, having a mean particle diameter of 30 μm and a bulk density of 0.3 g/ml.

Graphite F: flake, having a mean particle diameter of 10 μm and a bulk density of 0.23 g/ml.

Incidentally, the density of graphite was measured according to JIS K6891-1995.

*1: Resol-type phenolic resin, "PL-4804" from Gun-ei Kagaku Co., Ltd.

*2: Zinc stearate

[0053]

[Example 3] Polymer electrolyte fuel cell (1)

In this example, the unit cell is composed of a polymer electrolyte membrane ("Nafion") and a pair of electrodes (carbon paper from Chemics Co., Ltd.) holding the membrane between them. The membrane and electrodes were unified in the usual way. The unified electrode was held between the two separators prepared in Example 1. In this way there was obtained a unit cell having passages for fuel gas supply and discharge. Fifty pieces of the unit cells were assembled into a fuel cell by tightening with bolts and nuts. The separators did not crack at the time of assembling.

[0054]

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The resulting fuel cell was found to function satisfactorily.

It was tested for vibration and shock resistance in anticipation of their use on a car. The separators remained intact after 1000 cycles of vibration and shock.

[0055]

[Example 4] Polymer electrolyte fuel cell (2)

In this example, the unit cell is composed of a polymer electrolyte membrane ("Nafion") and a pair of electrodes (carbon paper from Chemics Co., Ltd.) holding the membrane between them. The membrane and electrodes were unified in the usual way. The unified electrode was held between the two separators prepared in Example 2. In this way there was obtained a unit cell having passages for fuel gas supply and discharge. One hundred pieces of the unit cells were assembled into a fuel cell by tightening with bolts and nuts. The separators did not crack at the time of assembling.

[0056]

The resulting fuel cell was found to function satisfactorily.

It was tested for vibration and shock resistance in anticipation of their use on a car. The separators remained intact after 1000 cycles of vibration and shock.

[BRIEF DESCRIPTION OF THE DRAWING]

5 [Fig. 1]

Fig. 1 is a perspective view showing one example of the fuel cell.

[Fig. 2]

Fig. 2 is a perspective view showing the fuel cell separator pertaining to one embodiment of the present invention.

[Fig. 3]

Fig. 3 is a schematic diagram showing the vertical cross section of the separator.

15 [Explanation of Symbols]

- 1 Separator
- la Rib
- 2 Electrolytic membrane of solid high polymer
- 3 Gas diffusion electrode
- 20 4 Gas feed groove (passage)

2000-204728 [ABSTRACT]

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[Means for Solving the Problem]

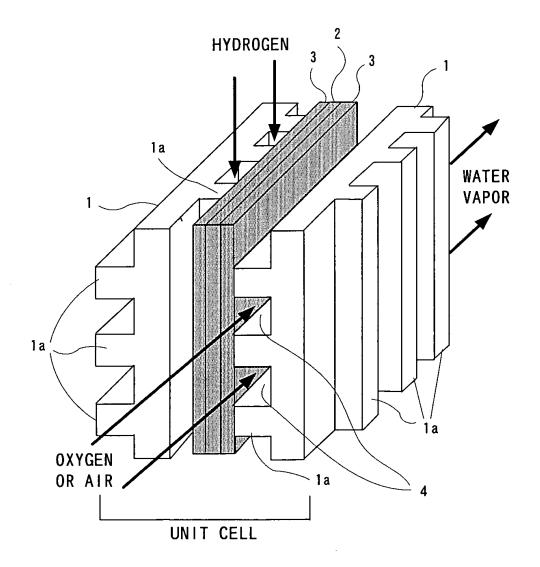
Disclosed herein are a fuel cell separator having gas supply grooves on one side or both sides thereof which is molded from a composition composed mainly of an electrically conductive carbon powder and a binding agent, wherein the electrically conductive carbon powder is present such that its particles longer than 70 μm at maximum in the major axis direction and longer than 30 μm at maximum in the minor axis direction along the vertical cross section of the fuel cell separator occupy more than 50% of the sectional area in the vertical direction, a process for production of the separator, and a polymer electrolyte fuel cell.

[Effect]

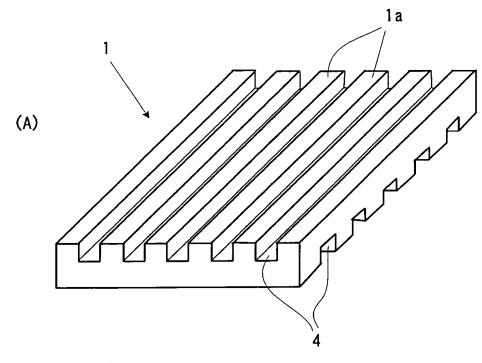
The present invention permits efficient mass production of high-quality fuel cell separators having high elasticity, high electrical conductivity, and good gas impermeability. The polymer electrolyte fuel cell in which all or part of the separators are those which pertain to the present invention exhibits good gas seal performance and good impact resistance without possibility of separators cracking at the time of assembling.

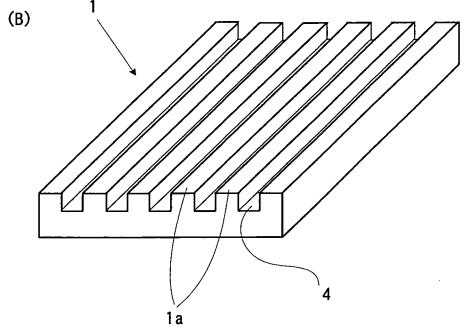
[Selected Drawing] FIG. 1

2000-204728 [DRAWING] [FIG. 1]

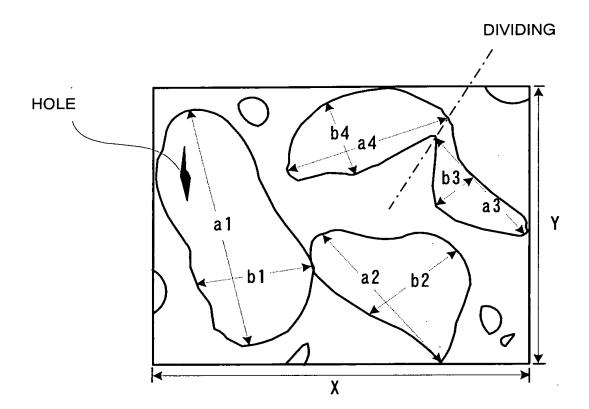


[FIG. 2]





[FIG. 3]



2000-204728

[Paper] APPLICATION FOR PATENT

[Reference Number] 12324

[Application Date] July 6, 2000

[Destination] The Commissioner of the Patent Office

Kozo OIKAWA

[International Patent Classification]

H01M 8/00

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